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[54] **DETERMINED DNA SEQUENCES DERIVED FROM A PAPILOMAVIRUS GENOME, THEIR USES FOR IN VITRO DIAGNOSTIC PURPOSES AND THE PRODUCTION OF ANTIGENIC COMPOSITIONS**

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[21] Appl. No.: **466,711**

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[52] U.S. Cl. **530/324**; 530/387.9; 530/350; 435/5; 435/235.1; 424/130.1; 424/186.1; 424/184.1
[58] Field of Search 435/69.3, 5, 235.1; 530/387.9, 389.4, 350, 324; 424/130.1, 186.1, 184.1

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[57] ABSTRACT

The invention concerns DNA fragments derived from the genomic DNA of HPV-33. These fragments are selected from the group of fragments extending between the nucleotide extremities defined hereafter in relation to the nucleotide-numbering in FIGS. 1a and 1b respectively:

- 76-556
- 543-864
- 867-2811
- 2728-3808
- 3326-3575
- 3842-4079
- 4198-5611
- 5516-8091.

The invention also relates to the use of these fragments as probes for the detection of HPV in tissue cultures.

8 Claims, 3 Drawing Sheets

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1 GTAAACTATA ATGCCAAGTT
 2 GCACGACTAT GTTTCAGGAC
 3 GGAATGCAAA AAAECTTTGC
 4 CTGTGTTTGC AATATTAAT
 5 TTTGGCAGGG AATATTAAT
 6 AATATGTTTT AATATTAAT
 7 ACAAGCACAA AATATTAAT
 8 CGAACCATAC AATATTAAT
 9 ATGGGGCTGG AATATTAAT
 10 TGACAGTGGC AATATTAAT
 11 GGGAGGATG AATATTAAT
 12 GACAGAGTGC AATATTAAT
 13 ATGTAGATA AATATTAAT
 14 ATGGATAAG AATATTAAT
 15 AGAAGCTTAA AATATTAAT
 16 AGGTGAGCA AATATTAAT
 17 AAACAAGTTC AATATTAAT
 18 AATGATGTC AATATTAAT
 19 GATGATGTC AATATTAAT
 20 GATGATGTC AATATTAAT
 21 GATGATGTC AATATTAAT
 22 GATGATGTC AATATTAAT
 23 GATGATGTC AATATTAAT
 24 GATGATGTC AATATTAAT
 25 GATGATGTC AATATTAAT
 26 GATGATGTC AATATTAAT
 27 GATGATGTC AATATTAAT
 28 GATGATGTC AATATTAAT
 29 GATGATGTC AATATTAAT
 30 GATGATGTC AATATTAAT
 31 GATGATGTC AATATTAAT
 32 GATGATGTC AATATTAAT
 33 GATGATGTC AATATTAAT
 34 GATGATGTC AATATTAAT
 35 GATGATGTC AATATTAAT
 36 GATGATGTC AATATTAAT
 37 GATGATGTC AATATTAAT
 38 GATGATGTC AATATTAAT
 39 GATGATGTC AATATTAAT
 40 GATGATGTC AATATTAAT
 41 GATGATGTC AATATTAAT
 42 GATGATGTC AATATTAAT
 43 GATGATGTC AATATTAAT
 44 GATGATGTC AATATTAAT
 45 GATGATGTC AATATTAAT
 46 GATGATGTC AATATTAAT
 47 GATGATGTC AATATTAAT
 48 GATGATGTC AATATTAAT
 49 GATGATGTC AATATTAAT
 50 GATGATGTC AATATTAAT
 51 GATGATGTC AATATTAAT
 52 GATGATGTC AATATTAAT
 53 GATGATGTC AATATTAAT
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 60 GATGATGTC AATATTAAT
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 63 GATGATGTC AATATTAAT
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 66 GATGATGTC AATATTAAT
 67 GATGATGTC AATATTAAT
 68 GATGATGTC AATATTAAT
 69 GATGATGTC AATATTAAT
 70 GATGATGTC AATATTAAT
 71 GATGATGTC AATATTAAT
 72 GATGATGTC AATATTAAT
 73 GATGATGTC AATATTAAT
 74 GATGATGTC AATATTAAT
 75 GATGATGTC AATATTAAT
 76 GATGATGTC AATATTAAT
 77 GATGATGTC AATATTAAT
 78 GATGATGTC AATATTAAT
 79 GATGATGTC AATATTAAT
 80 GATGATGTC AATATTAAT
 81 GATGATGTC AATATTAAT
 82 GATGATGTC AATATTAAT
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 84 GATGATGTC AATATTAAT
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 86 GATGATGTC AATATTAAT
 87 GATGATGTC AATATTAAT
 88 GATGATGTC AATATTAAT
 89 GATGATGTC AATATTAAT
 90 GATGATGTC AATATTAAT
 91 GATGATGTC AATATTAAT
 92 GATGATGTC AATATTAAT
 93 GATGATGTC AATATTAAT
 94 GATGATGTC AATATTAAT
 95 GATGATGTC AATATTAAT
 96 GATGATGTC AATATTAAT
 97 GATGATGTC AATATTAAT
 98 GATGATGTC AATATTAAT
 99 GATGATGTC AATATTAAT
 100 GATGATGTC AATATTAAT

FIG.1A

3701 ATGGAAATTGT AACGTGTAACA TTGTAACTG AACAGCAACA ACAAA TGT TTAGGTACC G TAAANAATACC ACCTACTGTG CAAATAAGTA CIGGATTTAT
 3801 GACATATATA CAAGCCAATA CAAGCCACTA TGCTGCTCTA ATTGATATA ACCAIGATAT ATATTGTTTT TATGCTTATC
 3901 CTATATATA TACTTCCAT TCTACTGAT CCAATGATGT TCTACTGAT GCTTGGTGGT TGGGTGTTTG TGGGTGTTTG
 4001 TTTTTTGT ATTTGTTGTT TTATATTTA TTATATTTA TTTATATTTA TTTATATTTA TTTATATTTA TTTATATTTA
 4101 TGTITGATA TAGTGCACA TGGTGGTGT TAAACATGT TAAACATGT TAAACATGT TAAACATGT TAAACATGT
 4201 TGTATATTA TGAGACAAA ACCATCTACA AGCCGCAAGC GTCATCTGC GTCATCTGC GTCATCTGC GTCATCTGC
 4301 TTATCTAA AGTGGAAAGCA AGTACCATAG CAGATCAAA TCTTAAATAT TCTTAAATAT TCTTAAATAT TCTTAAATAT
 4401 TTCAAGTGGG AGGACTGACT ATGTAACCTA ATGTAACCTA ATGTAACCTA ATGTAACCTA ATGTAACCTA ATGTAACCTA
 4501 GACAGTTAG ACTCCGCTAT AGTGTATAT ATGAGAGAAA CAAGTTTTAT AGAGGAGGT GAGAGGAGGT GAGAGGAGGT
 4601 TTACTACATC TCCAGATAGT ACACTGCAA TTATTAATGT TTATTAATGT TTATTAATGT TTATTAATGT TTATTAATGT
 4701 TGAACCATCT GTACTACACC CTCCAGGCC CTCCAGGCC CTCCAGGCC CTCCAGGCC CTCCAGGCC CTCCAGGCC
 4801 ATGGATACCT TTGTGTTTC CACAGACACT AGTATGTA CACAGACACT CACAGACACT CACAGACACT CACAGACACT
 4901 ATACCAACA GTTAAAGT TTGATAGCT GATGATATC ACTGCTCTT ACTGCTCTT ACTGCTCTT ACTGCTCTT
 5001 CACATACAAA TTTAAACA TTGATAGCT GATGATATC ACTGCTCTT ACTGCTCTT ACTGCTCTT ACTGCTCTT
 5101 ACTGTGGT TTAGTAAAGT AGGTCAAA GGCACACTTA AAATCGCAG TTACATGATA TTACATGATA TTACATGATA
 5201 TTGTGCTT AGACDACC CTGCCCAAATG CTGCCCAAATG CTGCCCAAATG CTGCCCAAATG CTGCCCAAATG
 5301 TGTATATCT CACGATGTTG ATAAATACA CACCCCAATG CACCCCAATG CACCCCAATG CACCCCAATG CACCCCAATG
 5401 ACAGATTTG ATACTCTCTG TTGCTGCTG CTGCTGCTG CTGCTGCTG CTGCTGCTG CTGCTGCTG CTGCTGCTG
 5501 ACAACATGT TGTAGACGCT GCTGACTTG TTATACATC TTATACATC TTATACATC TTATACATC TTATACATC
 5601 TGTGGGCC TTGTCAGGCC ACAGTGTACC TGTCTCTGT ACCTGTACT ACCTGTACT ACCTGTACT ACCTGTACT
 5701 TGTCTGAGT TCCAGACTC TTGCTGTTGG CCACTCATAT TTCTTATA AAATCGTAC AAATCGTAC AAATCGTAC
 5801 CAATATAAG TTTTAGGT CCGTTAGCA GATCCTAATA AAATGCTT TTTAGGCTT TTTAGGCTT TTTAGGCTT
 5901 TGTAGGCT TGAATAGGT AGAGGCGAG CATTAGGCT TTTAGGCT TTTAGGCT TTTAGGCT TTTAGGCT
 6001 TCTTGACAAA CCGGTGCTG TACTAATGCA GDCCTGCA ATGTTATCC ATGTTATCC ATGTTATCC ATGTTATCC
 6101 TGGGTAAAG GTTGTCTTG TACTAATGCA GDCCTGCA ATGTTATCC ATGTTATCC ATGTTATCC ATGTTATCC
 6201 CAGGATTTG ATGACTGAT TTTAAACAT TTTAAACAT TTTAAACAT TTTAAACAT TTTAAACAT TTTAAACAT
 6301 AATGACTGT GAGCTTATG GTGATAGT ATTTCTT TTTAAACAT TTTAAACAT TTTAAACAT TTTAAACAT
 6401 GCTTCTCG ATGACTGAT TTTAAACAT TTTAAACAT TTTAAACAT TTTAAACAT TTTAAACAT TTTAAACAT
 6501 AALCTAGT ALLAAAG CATA TTTG CATA TTTG CATA TTTG CATA TTTG CATA TTTG CATA TTTG CATA
 6601 CALYCCAGT ACTAATGCA CATA TTTG CATA TTTG CATA TTTG CATA TTTG CATA TTTG CATA TTTG CATA
 6701 GALTACAGT TGTCTTCA ACTATGCAA ACTATGCAA ACTATGCAA ACTATGCAA ACTATGCAA ACTATGCAA
 6801 TGGTTAGC ACTCTTCA ACTATGCAA ACTATGCAA ACTATGCAA ACTATGCAA ACTATGCAA ACTATGCAA
 6901 GAAAGATCC TTAGGTAAT ATAACTTCA TTAGGTAAT ATAACTTCA TTAGGTAAT ATAACTTCA TTAGGTAAT
 7001 CAGGACGCT TTAGGTAAT ATAACTTCA TTAGGTAAT ATAACTTCA TTAGGTAAT ATAACTTCA TTAGGTAAT
 7101 TGAATAGT TTAGGTAAT ATAACTTCA TTAGGTAAT ATAACTTCA TTAGGTAAT ATAACTTCA TTAGGTAAT
 7201 GTGATAGT ACTGCTTG TTTATGCT ACTGCTTG TTTATGCT ACTGCTTG TTTATGCT ACTGCTTG
 7301 GTATAGT TAACTATG ACTATGTTA TTTATGCT ACTGCTTG TTTATGCT ACTGCTTG TTTATGCT ACTGCTTG
 7401 CTTATTTCC CATA TTTG ACTATGTTA TTTATGCT ACTGCTTG TTTATGCT ACTGCTTG TTTATGCT ACTGCTTG
 7501 AACTGTTCC GTTATTTG CATACATACC CTTATTTG CATACATACC CTTATTTG CATACATACC CTTATTTG
 7601 ATACATACC TATGACATTT GACAGACAGT TATGACATTT GACAGACAGT TATGACATTT GACAGACAGT
 7701 TGCMAATA CTATATGTA CTAAATGTT ACACATGCT TATGACATTT GACAGACAGT TATGACATTT
 7801 CTTTGTATGC CAAACTATGC CTTGTAAG CTTTGTATGTT TATGACATTT GACAGACAGT TATGACATTT
 7901 TATATAATA

FIG.1B

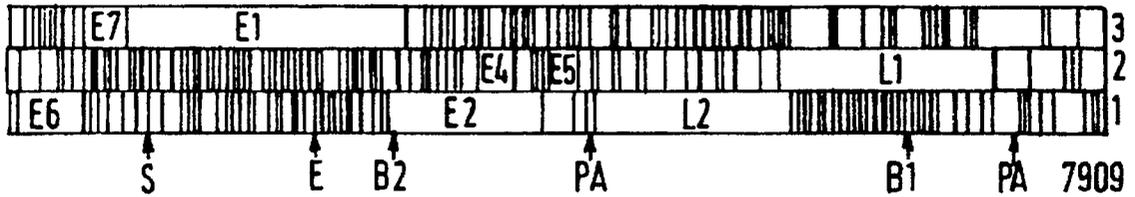


FIG.2

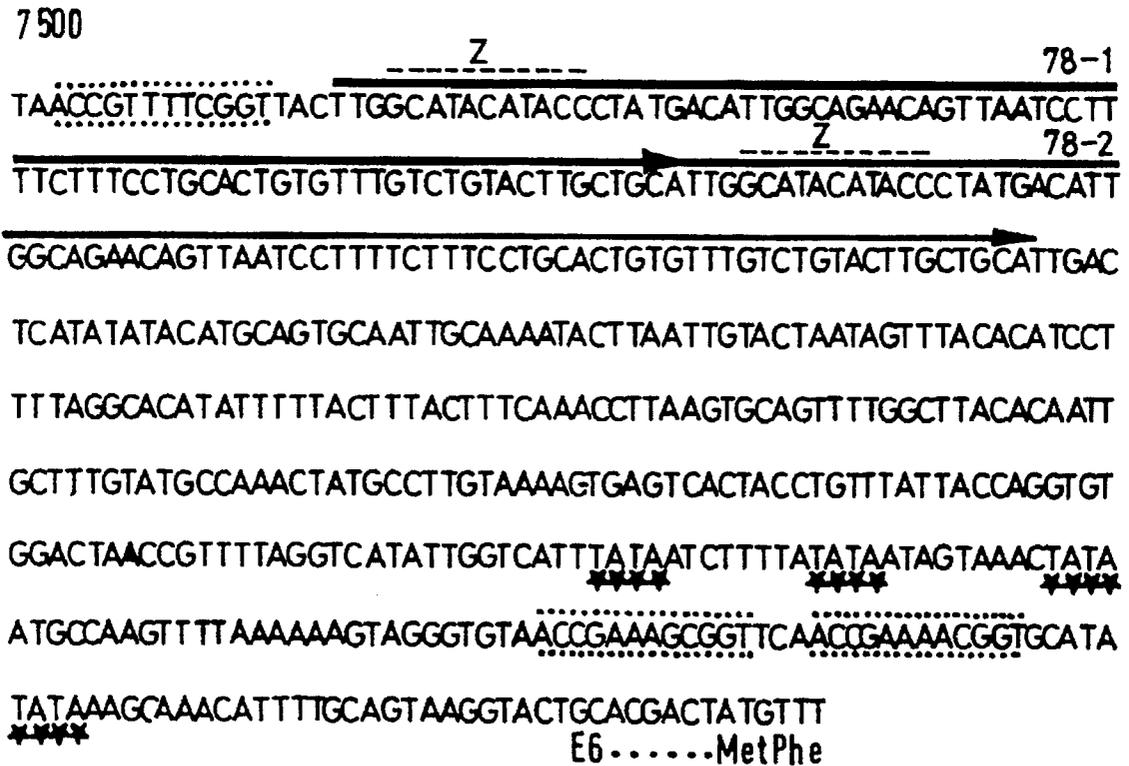


FIG.3

**DETERMINED DNA SEQUENCES DERIVED
FROM A PAPILLOMAVIRUS GENOME,
THEIR USES FOR IN VITRO DIAGNOSTIC
PURPOSES AND THE PRODUCTION OF
ANTIGENIC COMPOSITIONS**

This is a division of application Ser. No. 08/222,569, filed Mar. 25, 1994; U.S. Pat. No. 5,554,538; which is a continuation of application Ser. No. 08/161,239 filed Nov. 10, 1993; abandoned; which is a continuation of application Ser. No. 08/032,694 filed Mar. 17, 1993; abandoned; which is a continuation of application Ser. No. 07/908,895 filed Jul. 8, 1992; abandoned; which is a continuation of application Ser. No. 07/664,503 filed Mar. 5, 1991; abandoned; which is a continuation of application Ser. No. 07/518,302 filed May 2, 1990; abandoned; which is a continuation of application Ser. No. 07/128,341, filed as PCT/EP87/00158, Mar. 20, 1987 abandoned.

The invention pertains to determined DNA sequences derived from a papillomavirus genome, more particularly DNA recombinants, including vectors, modified by such DNA sequences in such manner that, when said DNA recombinants are introduced in suitable host cells in which said DNA recombinants can be replicated, the said DNA sequences can be expressed in the form of the corresponding proteins. The invention further relates to the proteins themselves, which can be purified and used for the production of immunogenic compositions.

The invention pertains more particularly to DNA products of the papillomavirus designated as IP-2 (now redesignated as HPV-33) in the European patent application filed under number 85.402362.9 on Nov. 29, 1985, the contents of which are incorporated herein by reference. A plasmid containing the DNA of said virus has been deposited at the CNCM ("Collection nationale de Culture de Micro-Organismes" of the Pasteur Institute of Paris) under number I-450.

Papillomaviruses are members of the papovavirus family and possess a genome of about 7,900 base pairs (bp) consisting of a covalently closed circular DNA molecule. Human papilloma viruses (HPV) are classified on the basis of their DNA sequence homology (6) and nearly 40 types have now been described. Considerable insight into HPV biology and their involvement in human disease has been attained by the application of the techniques of molecular biology. A possible role for HPVs in human cancer was suspected following the detection of HPV DNA in tumors resulting from the malignant conversion of genital warts (33). The cloning of two HPV genomes, HPV-16 and HPV-18 (3, 11) from cervical carcinomas has further stimulated research in this field of immense socio-economic importance. These viruses were discovered in more than 70% of the malignant genital tumors examined and in many others HPV-16 related sequences were detected (3, 16, 33). Amongst these is HPV-33 which was recently cloned from an invasive cervical carcinoma using HPV-16 as a probe under conditions of reduced stringency (1). In the present study we have determined the DNA sequence of HPV-33 and describe its relationship to HPV-16. Among the papillomaviruses HPV-33 is unique as it possesses a 78 bp tandem repeat which strongly resembles the enhancer of SV40 (4, 14).

The invention stems from the cloning strategy disclosed hereafter of the genome of HPV-33 which enabled particular DNA sequences to be identified, more particularly those providing hybridization probes, particularly useful for the detection of DNA of papillomaviruses related to HPV-33 in

human tissue, whereby positive responses can be related to the possible development in the host of invasive cervical carcinomas.

Reference is hereafter made to the drawings in which the 5 Figures concern respectively:

FIGS. 1a and 1b. Nucleotide sequence of HPV-33. Position 1 on the circular genome corresponds to a "Hpa-like" sequence found by alignment with HPV-6b.

FIG. 2. Distribution of the major reading frames in the HPV-33 genome. The reading frames were identified by comparison with other HPV sequences and the stop codons are represented as vertical bars. Also indicated are the locations of unique restriction sites (S, SmaI; E, EcoRV; B2, BglII; B1, BglI) and the likely polyadenylation signals (PA) for the early and late transcripts. In addition to these, 6 other potential PA sites (AATAAA) were detected at positions 862, 1215, 1221, 2666, 5837 and 6239.

FIG. 3. Principle features of the non-coding region. A section of the non-coding region from positions 7500 to 114 is shown. The 78 bp tandem repeats are overlined and those regions resembling the Z-DNA forming element of the SV-40 enhancer are indicated. Potential promoter elements are denoted by stars and the 3 copies of the 12 bp palindrome enclosed between two rows of dots.

Preferred sequences are those which encode full proteins, more particularly and respectively the nucleotidic sequences having the open reading frames referred to in table I hereafter.

The conditions under which the DNA sequence analysis were performed are defined under the heading "MATERIALS AND METHODS" hereafter. The conclusions which were drawn from this sequence analysis appear under the heading "DISCUSSION".

MATERIALS AND METHODS

DNA sequence analysis.

The source of HPV-33 sequenced in this study was plasmid p15-5 (1) which consists of a BglIII linearized HPV-33 genome cloned in a pBR322 derivative. A library of random DNA fragments (400-800 bp) was prepared in M13mp8 (17) after sonication and end-repair of p15-5, essentially as described previously (28). DNA sequencing was performed by the dideoxy chain termination method (19, 20) with the modifications of Biggin et al. (2). Most of the sequence was derived in this way although part of the non-coding region was found to be absent or under-represented in the M13 library (>300 clones). The sequence of this region was obtained directly from p15-5 using the method of Smith (24). Briefly, restriction fragments isolated from 2 "complementary" M13 clones were used to prime DNA synthesis on templates prepared from p15-5 which had been linearized with a restriction enzyme and then treated with exonuclease III (200 units/pmol DNA for 1 h at 22° C.).

Computer analysis.

DNA sequences were compiled and analysed with the programs of Staden (26, 27) as modified by B. Caudron. Optimal alignments of DNA or protein sequences were obtained using the algorithm developed by Wilbur and Lipman (31).

RESULTS AND DISCUSSION

Genomic Arrangement of HPV-33

The complete 7909 nucleotide sequence of HPV-33, determined by the M13 shotgun cloning/dideoxy sequencing approach, is presented in FIG. 1. On average each position was sequenced 6.5 times. In agreement with the convention

for other papillomavirus sequences the numbering begins at a site resembling the recognition sequence for HpaI in the non-coding region.

An analysis of the distribution of nonsense codons (FIG. 2) shows that, as in all other sequenced papillomaviruses, the 8 major open reading frames are located on the same strand. Some features common to HPV-33 and HPV types 1a, 6b and 16 together with the cottontail rabbit papillomavirus and the prototype bovine papillomavirus, BPV-1, (5, 7, 8, 13, 21, 22) include the overlap between the largest open reading frames in the early region, E1 and E2, and the inclusion of E4 within the section encoding E2. Interestingly, the BglII site used in the molecular cloning of HPV-33 is situated within the E1/E2 overlap. Another property common to all papillomaviruses, except BPV-1, is the overlap between the L1 and L2 reading frames. Following L1 is the 892 bp non-coding region which, by analogy with BPV1 (15, 29) undoubtedly contains the origin of replication and various transcriptional regulatory elements. The principal characteristics of the HPV-33 genome are summarized in Table 1.

Nucleotide Sequence Comparison with HPV-16

HPV-16 is the only other oncogenic papillomavirus, isolated from tumors of the ano-genital region, which has been completely sequenced (22). The gross features of HPV-33 resemble those of HPV-16 except that the E1 reading frame of the latter is interrupted. All of the coding sequences in HPV-33, except that of E5, are slightly shorter than their counterparts in HPV-16. This may contribute to the fact that its non-coding region, between L1 and E6 (FIG. 2), is 76 bp longer thereby keeping the genomes nearly constant in size.

When the open reading frames were compared pairwise (Table 2) it was found that E1, E2, E6, E7, L1 and L2 displayed between 65–75% homology whereas those for E4 and E5 were more divergent (about 50% homology). These findings confirm the heteroduplex analysis performed previously (1). A comparative study (8) of papillomavirus E1 gene products showed that the polypeptide consists of an NH₂-terminal segment whose sequence is highly variable, and a COOH-terminal domain of well-conserved primary structure. The longest stretch of perfect sequence homology, 33 nucleotides (positions 1275–1307, FIG. 1) is found near the 5'-end of the E1 reading frame in a region encoding the variable domain of the polypeptide. Several other regions of complete identity (19–28 nucleotides) were detected elsewhere in E1, and also in E2, L2 and as many of these sequences are not found in the genomes of other HPVs, such as HPV-1a and HPV-6b, this raises the possibility that the corresponding oligonucleotides could be produced and used as diagnostic hybridization probes for screening biopsy material from potentially tumorigenic lesions.

Potential Gene Products

The papillomavirus gene products may be divided into those which are believed to play a purely structural role, L1 and L2, and those required for viral propagation and persistence. The results of a comparison of the probable products of the major reading frames from HPVs-33, 16 and 6b are summarized in Table 2. As expected there is strong identity between the oncogenic HPVs-33 and 16, particularly for the proposed E1, E6, E7, L2 and L1 proteins. When conservative substitutions are included the homology between the two L1 polypeptides increases to 90% suggesting that the corresponding capsids must be antigenically related. In contrast, significantly weaker homologies were detected when the analysis was extended to include the benign genital wart-forming HPV-6b (Table 2). Comparison

of the HPV-16 proteins with those of HPV-6b revealed slightly more homology than was found with HPV-33 suggesting a closer evolutionary relationship.

The non-coding Region

The non-coding region of HPV-33 displays several unique properties and bears only weak resemblance to its homologue in HPV-16. Located between the L1 stop codon and including the putative polyadenylation signal for the late transcripts is a stretch of 223 bp (positions 7097–7320, FIG. 1) unusually rich in T+G (79%). Contained within this segment are two copies of a 19 bp direct repeat (with one mismatch) and 7 copies of the motif TTGTRTR (where R is A or G). The latter is also found 7 times in the corresponding region of HPV-16 suggesting that it may represent a recognition site for proteins involved in replication. It should be noted that nascent replication forks have been localised in this region of the BPV-1 genome (29) and that the origin of replication of the Epstein-Barr virus consists of a family of repeated sequences (32).

A 12 bp palindrome (ACCG . . . CCGT) that occurs exclusively in the non-coding region of all papillomavirus genomes examined was recently reported by Dartmann et al. (9). Three copies were found in the HPV-33 genome (FIG. 3) and these occupy the same positions in the non-coding region of HPV-16. A role for the palindrome as a possible control site for the early promoter was proposed (4, 9, 15) and indirect support is provided by our finding that the non-coding regions of HPVs, such as HPV-33, do not display the clustered arrangement of recognition sites for the promoter-specific, activation factor Sp1(12). This is in direct contrast to the situation in another papovavirus, SV40 (12, 14).

The most striking feature of HPV-33 is a perfect 78 bp tandem repeat located 200 bp after the putative origin of replication (FIG. 3). No other repeats of this size or sequence have been described in the genomes of other papillomaviruses. The presumed early promoter for HPV-33 is located about 300 bp downstream from the tandem repeat and the characteristic promoter elements (4) could be identified (FIG. 3). The size, position and arrangement of the 78 bp repeats in the HPV-33 genome suggest that they may function as enhancers of viral transcription. Tandem repeats of 72, 73 and 68 bp have been located near the early promoter of SV40 (4, 14), in the LTR of moloney murine sarcoma virus (10), and in the BK virus genome (23) and shown to enhance transcription from PolII dependent promoters in a cis-active manner. From mutagenesis of the SV40 enhancer (14, 30) and sequence comparisons of characterized transcriptional activators a consensus enhancer sequence was derived. This structure could not be detected in the 78 bp repeat but a potential Z-DNA forming region was uncovered. Z-DNA is believed to attract regulatory molecules to eukaryotic promoters and a Z-DNA antibody binding site has been demonstrated within the SV40 enhancer (18). The sequence to which this antibody binds is also found, albeit with a single mismatch, in the putative HPV-33 enhancer (positions 7520–7527, 7599–7606, FIGS. 1, 3).

The proposed HPV-33 enhancer shows no extended sequence homology to the well-characterized enhancers nor to other papillomavirus regulatory regions. However, it has recently been demonstrated that an enhancer-like element is located in the non-coding region of BPV-1 and that it requires the E2 product for activation (25). These findings support our proposal that the 78 bp tandem repeats could have enhancer function and may indicate that the relatively low homology (Table 2) between the E2 proteins of HPV-33

and 16 reflects a specificity for the corresponding enhancer/regulatory regions.

Tables 1 and 2 which have been referred to in the instant disclosure follow.

TABLE 1

Principal features of the HPV-33 genome				
Open Reading Frame	START	FIRST ATG	STOP CODON	mol.wt.
E6	76	109	556 TGA	17 632
E7	543	573	854 TAA	10 825
E1	867	879	2811 TGA	72 387
E2	2728	2749	3808 TAA	40 207
E4	3326	—	3575 TAG	9 452
E5	3842	—	4079 TAA	9 385
L2	4198	4210	5161 TAG	50 539
L1	5516	5594	7091 TAA	55 839

a. Calculated from the first ATG where this exists or from the start of the open reading frame.

TABLE 2

Comparison of HPV proteins ^a			
Protein	HPVs		
	33v16	33v6b	16v6b
E6	65(70)	36(51)	37
E7	61(69)	55(60)	56
E1	61(69)	50(60)	53
E2	53(65)	46(58)	45
E4	52(55)	39(46)	48
E5	40(52)	39(43)	33
L2	64(66)	52(58)	53
L1	81(75)	68(69)	71

^aExpressed as % homology after alignment with the program of (31). Values in parenthesis represent % nucleotide sequence homology.

The invention relates more particularly to sequences corresponding to the open reading frames of E6, E7, E1, E2, E4, E5, L2, L1.

The invention pertains also the uses of these sequences as hybridization probes, either those which are useful also for the detection of other papillomaviruses, thus of groups of papillomaviruses—such as probes containing part or all of the open reading frames corresponding to L1—or those which are more virus—specific, i.e. probes containing part or all of the open reading frame corresponding to.

It also relates to other probes which detect sub-groups of papillomaviruses, particularly probes for the detection of viruses which can be related to major classes of diseases, i.e. viruses associated with tumors. By way of example of one of said probes one should mention that which contains the sequence positioned between nucleotides 1275 and 1307 according to the numbering of the nucleotides in FIGS. 1A, 1B.

Needless to say that the invention also pertains to all of said DNA sequences, when labelled by a suitable label, i.e. a radioactive enzymatic or immunofluorescent label.

DNAs derived from the viral genome and which carry nucleotides modified by a chemical group which can be recognized by antibodies also form part of the invention. It is well known that such DNAs can be produced by nick-translation in the presence of nucleotides modified accordingly. These DNAs form particularly valuable hybridization probes which, when hybridized to a DNA preparation containing the complementary strand sought, can be detected by the above mentioned antibodies.

The invention also pertains to the diagnostic methods *per se*. Suitable methods are exemplified hereafter.

Several hybridization methods may be used. For example, the spot hybridization method includes, after denaturation of the DNA, the deposition of an aliquot of the DNA onto film supports (nitrocellulose or Genescreenplus), the hybridization of each film under the usual conditions with the probe, and the detection of the radioactive hybrid by contact exposition of the hybridized film onto radiographic film. Another possibility is replicated culture hybridization which involves agarose gel electrophoresis separation of the DNA fragments resulting from treatment of the DNA by restriction enzymes, the transfer of the fragments after alkaline denaturation onto films (nitrocellulose or Genescreenplus) and their hybridization under usual conditions with different mixtures of probes. The formation of radioactive hybrids is detected again by contact exposition of the hybridization support films onto radiographic film.

For instance the probes of the invention can be used for the detection of the relevant viruses (or DNAs thereof) in preparation consisting of a biopsy of cells obtained by scraping a lesion, or of biopsy sections fixed with Carnoy's mixture (ethanol, chloroform, acetic acid 6:3:1) and included in paraffin.

The above nucleotide sequences can be inserted in vectors, to provide modified vectors which, when introduced in the suitable cell host, are capable of providing for the transcription and, where appropriate, translation of said DNA sequences to produce the corresponding proteins which can then be isolated from cellular extracts of the hosts. Obviously it is within the knowledge of the man skilled in the art to select the appropriate vectors, particularly in relation to the host to be transformed therewith. Vectors consist for instance of plasmids or phages which will be selected according to their recognized capability of replicating in the corresponding prokaryotic cells (or yeast cells) and of allowing for the expression of the DNA sequence which they carry.

The invention also relates to DNA recombinants containing an insert consisting of a DNA sequence corresponding to any of the above-defined open reading frames or of a part thereof, and suitably engineered to allow for the expression of the insert in eucaryotic cells, particularly cells of warm-blooded animal. Suitable DNA recombinants are genetic constructs in which said insert has been placed under the control of a viral or eucaryotic promoter recognized by the polymerases of the selected cells and which further comprise suitable polyadenylation sites downstream of said insert.

By way of example, the invention pertains to DNA recombinants containing any of the above-mentioned open-reading inserts placed under the control of a promoter derived from the genome of the SV40 virus. Such DNA recombinants—or vectors—can be used for the transformation of higher eucaryotic cells, particularly cells of mammals (for instance Veto cells). The invention further pertains to portions of the above identified DNA sequences which, when inserted in similar vectors, are able to code for portions of the corresponding proteins which have immunological properties similar to those encoded by the full nucleotide sequences mentioned above. The similarity of immunological properties can be recognized by the capacity of the corresponding polypeptides produced by the relevant host to be recognized by antibodies previously formed against the proteins produced by the cells previously transformed with vectors containing the above mentioned entire DNA sequences.

It goes without saying that the invention also pertains to any nucleotidic sequence related to the preceding ones which may be obtained at least in part synthetically, and in which the nucleotides may vary within the constraints of the genetic code, to the extent where these variations do not entail a substantial modification of the polypeptidic sequences encoded by the so-modified nucleotidic sequences.

It already flows from the preceding discussion that the invention also pertains to the purified proteins or polypeptides themselves as obtainable by the methods discussed hereabove. These polypeptides, when produced in a suitable host, can either be obtained from the cells, for instance after rupturing of their cell walls, or from the culture medium of said cells when excreted in said cell medium, depending on the cell DNA recombinant system which is used. The polypeptide obtained can then be purified by resorting to usual purification procedures. It should be understood that "purified" in the instant context means a level of purity such that, when electrophoresed in SDS-PAGE, the purified proteins yield a single detectable band, say by Western blot.

The viral proteins obtained, more particularly the structural proteins, for instance as a result of the expression of said DNA sequences in *E. coli*, can be used for the in vitro detection of antibodies against papillomavirus likely to be detected in tissue samples of patients possibly infected with papillomavirus.

Of particular relevance are the genetically engineered proteins having the peptidic sequences which can be deduced from the L1 and L2 open reading frames. Another peptide of interest is the E6* protein (E6 star), the synthesis of which can be induced by splicing and which encoded by a nucleotidic sequence located between nucleotides 229 (donor site) and 404 (acceptor site) of the HPV 33 sequence (see more particularly FIG. 1A), which sites also define the putative splicing sites in the E6* open reading frame of HPV 33. Reference may be had to the publication of Schneider-Gardicke and Schwartz, *Embo. J.*, 5, 2285-2292, as concerns the conditions of the production of such proteins.

These purified polypeptides can in turn be used for the production of corresponding antibodies which can be used for diagnosing in vitro the presence of viral polypeptides in a biological fluid, particularly in a serum or tissue culture of a patient. Like in the preceding instance, the invention relates to portions of the above defined polypeptides, particularly those which are recognized by the same antibodies or to the contrary are able to elicit in vivo the production of antibodies recognizing the complete proteins.

It must be understood that the inventions relates also specifically to the particular peptides encoded by the DNA regions specifically referred to in the preceding disclosure and which have been found of particular interest.

The invention further concerns host cells transformed with DNA recombinants containing nucleotidic sequences directing the expression of the different peptides mentioned hereabove, and effectively capable to produce said peptides when cultured in an appropriate culture medium.

The invention finally also pertains more particularly to the antibodies themselves which can be obtained from an animal, such as rabbit, immunized in standard manner with said purified polypeptides and/or from hybridomas previously prepared also in any known manner. Of particular interest are the antibodies (polyclonal and monoclonal antibodies) directed against the structural proteins. These antibodies are useful for the detection of viral infection. The antibodies which recognize the L1, L2 and E6* proteins of

HPV-33 are of particular significance. Antibodies specific of L2 provide diagnostic tools for the in vitro detection of specific viruses sharing with HPV-33 a sequence encoding a similar L2 protein. Antibodies specific to L1 are useful for the detection of the groups of viruses, to which HPV-33 belongs. Antibodies specific to the E6* protein are useful for the detection of the oncogenic character of the virus causing the abovesaid viral infection.

The invention also relates to intergenic sequences of particular interest, particular the 78 bp sequence. This sequence is of particular interest as a possible insert in eucaryotic vectors, particularly in a position upstream of the promoter and downstream of the site at which transcription of the gene or nucleotide sequence the transcription of which is sought is initiated in the relevant host.

All documents referred to herein are incorporated herein by reference. Particularly these documents can be referred to as concerns the definition of expressions used in this application where appropriate. As such they form part of the present disclosure.

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We claim:

1. A purified peptide obtained from a host cell transformed by a vector comprising a DNA fragment consisting of an open reading frame, which DNA fragment encodes said peptide and has a nucleotide sequence selected from the following group of nucleotide sequences numbered as shown in FIGS. 1a and 1b:

76-556
543-864
867-2811
2728-3808
3326-3575
3842-4079
4198-5611
5516-7091,

wherein said DNA fragment has been expressed.

2. A purified peptide obtained from a host cell transformed by a vector comprising a DNA fragment encoding said peptide, which DNA fragment has a nucleotide sequence selected from the following group of nucleotide sequences numbered as shown in FIGS. 1a and 1b:

109-556
573-864
879-2811
2749-3808
3326-3575
3842-4079
4210-5611
5594-7091,

wherein said DNA fragment has been expressed.

3. The purified peptide of claim 1, wherein said DNA fragment is cloned.

4. The purified peptide of claim 2, wherein said DNA fragment is cloned.

5. A purified antibody directed against the peptide of claim 1.

6. A purified antibody directed against the peptide of claim 2.

7. A purified antibody directed against the peptide of claim 3.

8. A purified antibody directed against the peptide of claim 4.

* * * * *

UNITED STATES PATENT AND TRADEMARK OFFICE
CERTIFICATE OF CORRECTION

PATENT NO. : 5,648,459
DATED : July 15, 1997
INVENTOR(S) : Cole et al.

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

In Table 1, column 5, line 17, the stop codon of open reading frame L2 should read --5611-- instead of "5161".

Signed and Sealed this
Fourteenth Day of April, 1998



Attest:

BRUCE LEHMAN

Attesting Officer

Commissioner of Patents and Trademarks